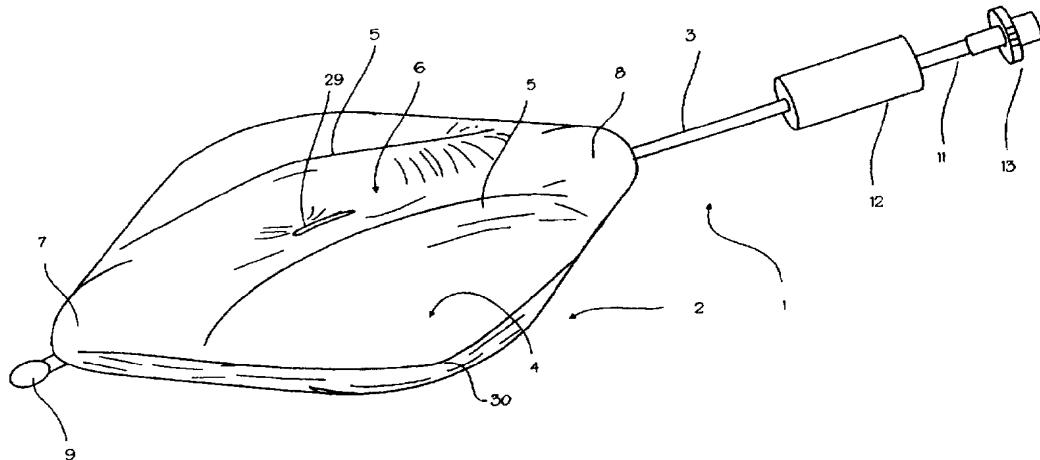


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(54) Title: MINIMALLY INVASIVE DIRECT CARDIAC MASSAGE DEVICE AND METHOD



(57) Abstract

This invention is a minimally invasive device for performing direct cardiac massage including an inflatable bladder (2) mounted on a rigid inflation tube (3). The rigid inflation tube is used to push the bladder into the sternocostal space through an incision in the upper abdomen just below the xiphoid process. After inserting into the sternocostal space, the bladder is repeatedly inflated and deflated to massage the heart and provide the blood flow.

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Minimally Invasive Direct Cardiac Massage Device And
Method

Field of the Invention

This invention relates to cardiopulmonary resuscitation and
5 cardiac massage.

Background of the Invention

Cardiopulmonary resuscitation (CPR) is a well known and valuable method of first aid. CPR is used to resuscitate people who have suffered from cardiac arrest after heart attack, 10 electric shock, chest injury and many other causes. During cardiac arrest, the heart stops pumping blood, and a person suffering cardiac arrest will soon suffer brain damage from lack of blood supply to the brain. Thus, CPR requires repetitive chest compression to squeeze the heart and the thoracic cavity 15 to pump blood through the body. Very often, the victim is not breathing, and mouth to mouth artificial respiration or a bag valve mask is used to supply air to the lungs while the chest compression pumps blood through the body.

It has been widely noted that CPR and chest compression can 20 save cardiac arrest victims, especially when applied immediately after cardiac arrest. Chest compression requires that the person providing chest compression repetitively push down on the sternum of the victim at 80-100 compressions per minute. CPR and closed chest compression can be used anywhere, wherever the 25 cardiac arrest victim is stricken. In the field away from the hospital, it may be accomplished by ill-trained by-standers or highly trained paramedics and ambulance personnel or at the hospital where it can be accomplished by doctors and nurses.

When a first aid provider performs chest compression well, blood flow in the body is typically about 25-30% of normal blood flow. This is enough blood flow to prevent brain damage. However, when chest compression is required for long periods of time, it is difficult if not impossible to maintain adequate compression of the heart and rib cage. Even experienced paramedics cannot maintain adequate chest compression for more than a few minutes. Hightower, et al., Decay In Quality Of Chest Compressions Over Time, 26 Ann. Emerg. Med. 300 (Sep. 1995).

10 Thus, long periods of CPR, when required, are not often successful at sustaining or reviving the victim. At the same time, it appears that, if chest compression could be adequately maintained, cardiac arrest victims could be sustained for extended periods of time. Occasional reports of extended CPR efforts (45-90 minutes) have been reported, with the victims eventually being saved by coronary bypass surgery. See Tovar, et al., Successful Myocardial Revascularization and Neurologic Recovery, 22 Texas Heart J. 271 (1995).

20 Mechanical devices for closed chest compression have been proposed and used. The device shown in Barkolow, Cardiopulmonary Resuscitator Massager Pad, U.S. Patent 4,570,615 (Feb. 18, 1986), the commercially available thumper device, and other such devices, provide continuous automatic closed chest compression. However, these devices are not clinically more successful than manual chest compression. See Taylor, et al., External Cardiac Compression, A Randomized Comparison of Mechanical and Manual Techniques, 240 JAMA 644 (Aug. 1978).

25 Active compression decompression using a device which also lifts the chest wall after compression has been proposed. Stiel, et al., The Ontario Trial of Active Compression and Decompression Cardiopulmonary Resuscitation for In-Hospital and Prehospital

Cardiac Arrest, 275 JAMA 1417 (1996) compared active compression-decompression with standard CPR, and found no significant improvement in survival or neurological outcome. A variety of other methods of increasing the effectiveness of CPR 5 have been proposed, including abdominal binding and anti-shock pants. These techniques are intended to block blood flow to the abdomen and legs, thus directing blood flow to the brain. Again, these techniques have not proven effective in boosting the survival rate of cardiac arrest victims.

10 Chest compression must be accomplished vigorously if it is to be effective. Very little of the effort exerted in chest compression actually compresses the heart and large arteries of the thorax and most of the effort goes into deforming the chest and rib cage. The force needed to provide effective chest 15 compression creates risk of other injuries. It is well known that placement of the hands over the sternum is required to avoid puncture of the heart during CPR. Numerous other injuries have been caused by chest compression. See Jones and Fletter, Complications After Cardiopulmonary Resuscitation, 12 AM. J. 20 Emerg. Med. 687 (Nov. 1994), which indicates that lacerations of the heart, coronary arteries, aortic aneurysm and rupture, fractured ribs, lung herniation, stomach and liver lacerations have been caused by CPR. Thus the risk of injury attendant to chest compression is high, and clearly may reduce the chances of 25 survival of the victim vis-a-vis a resuscitation technique that could avoid those injuries. Also, chest compression will be completely ineffective for very large or obese cardiac arrest victims because the chest cannot be compressed enough to cause blood flow.

In the hospital setting, when closed chest compression is ineffective, doctors have the option of using open chest compression (also referred to as cardiac massage, open cardiac massage, open resuscitation, etc.). To accomplish open chest compression, the doctors perform a thoracotomy and pull the rib cage apart to open the chest, cut the heart out of the pericardial sac and expose the heart, and then "massage" the heart by hand (they squeeze it like a bladder pump). Mechanical devices for squeezing the heart during open chest surgery have been proposed, such as Goetz, Heart Massage Apparatus, U.S. Patent 4,048,990, which provides a tulip shaped or basket-shaped bladder to surround the heart after the chest has been opened and the heart dissected from the pericardial sac. The grossly invasive thoracotomy procedure required for manual or mechanical open heart massage can only be accomplished in the hospital, and it carries its own risk of killing the cardiac arrest victim. Open chest cardiac massage is viewed as a last resort. See Blakeman, Open Cardiac Resuscitation, A Surgeons Viewpoint, 87 Postgraduate Med. 247 (Jan. 1990). However, it has the benefit of increased blood flow compared to closed chest compressions, about 50% of normal blood flow. Bartlett, et al., Comparative Study Of Three Methods Of Resuscitation: Closed Chest, Open Chest Manual And Direct Mechanical Ventricular Assistance, 13 Ann. Emerg. Med. 773 (1984).

Direct cardiac massage can be accomplished without open heart surgery. Buckman, et al., Direct Cardiac Massage Without Major Thoracotomy, 29 Resuscitation 237 (1995) shows a cardiac compression device which has a small plate mounted on a handle, like a potato masher or a toilet plunger. The device is inserted through the chest wall, through an incision between the ribs which is 7.5 cm long. The device is placed so that the small

plate is in contact with the left ventricle, then it is manually pushed against the heart to squeeze the heart. In both open cardiac massage and Buckman's minimally invasive direct cardiac massage, blood flow is accomplished by mechanically squeezing 5 the heart so that it acts like a bladder pump.

Summary of the Invention

The cardiac pumping device and method presented below allows for direct cardiac massage in a procedure that can easily be accomplished by emergency medical personnel, paramedics, 10 doctors and nurses, and probably by anyone trained in first aid. The substernal cardiac pump includes an inflatable balloon mounted on a rigid tube. The tube has inflation ports opening into the balloon, and a long handle section which is attached to an air pump. The air pump is preferably a positive placement 15 dual action pump, so that it pumps air into the balloon on the pumping stroke and sucks air out of the balloon during the reset stroke. Thus the balloon can be repeatedly inflated and deflated. When inflated, the balloon has a shape which accommodates the heart and squeezes the heart.

20 To use the cardiac pump on a cardiac arrest victim, the medic makes a small incision (two or three centimeter is sufficient) just below the sternum. The incision is shallow, just enough to puncture the skin and any fat beneath the skin. After making this initial incision, the medic sticks his finger 25 through the incision, slides his finger along the under-surface of the sternum and pushes a hole up through the diaphragm of the victim. This creates a channel into the thorax of the victim. This is all the preparation that is needed for insertion of the cardiac pump. With the channel easily made, the medic pushes the

balloon through the channel, into the thorax, and in place over the heart. The medic then operates the air pump to inflate and deflate the balloon repeatedly. Every time the balloon is inflated, it expands between the sternum and the heart, and thus 5 squeezes the heart.

Various additional features are added to make the cardiac pump easy to use. The hand pump may be hooked up to a small electric motor, powered by battery, standard household current, or through an automobile cigarette lighter or other mobile power 10 sources carried by ambulances. A variety of other air pump mechanisms can be used. The cardiac pump can be provided with over-pressure relief valves to limit the pressure within the balloon. The cardiac pump can be provided with an low pressure warning system to indicate rupture of the balloon or a leak in 15 the inflation pathway. The size of balloon inflation can be adjusted up or down in response to feedback from information gathered from measurement of the air exhaled by the victim.

The cardiac pump makes it easy to provide adequate heart compression for extended periods of time, without loss of 20 effectiveness due to fatigue of medics. The placement of the cardiac pump is simply accomplished, and direct cardiac massage is accomplished without thoracotomy or large incision. Actual compression of the heart is limited by the size of the balloon 5 and/or the pressure limits of the inflation system.

25

Brief Description of The Drawings

Figures 1 and 2 show the cardiac pump.

Figures 3 and 4 show the cardiac pump inserted inside an introducer sheath.

Figure 5 is a detail of the inflation tube.

Figures 6 and 7 show details of the balloon

Figures 8 and 9 show cross sections of the balloon.

Figure 10 shows an exploded view of the pieces used to
5 fabricate the balloon of Figure 2.

Figure 11 shows an anterior view of a human chest and the
insertion pathway for the cardiac pump.

Figure 12 shows cross section of a human chest and
preparation of the insertion pathway for the cardiac pump.

10 Figure 13 shows cross section of a human chest and the 20
initial placement of the cardiac pump.

Figure 14 is an illustration of the cardiac pump placed
inside the chest of a human being and inflated to compress the
heart.

15 Figure 15 is an illustration of the cardiac pump deflated
25 inside the chest to allow expansion of the heart.

Figure 16 shows an anterior view of the human chest and the
position of the deployed cardiac pump.

Figure 17 is a schematic of a manual pumping mechanism for
20 the cardiac pump.

Detailed Description of the Invention

Figure 1 shows the cardiac pump 1. The balloon 2 is mounted
on the rigid inflation tube 3. The balloon has a rounded diamond
shape, or a manta ray shape when viewed from above. The

posterior surface 4 of the balloon has a railed shape when inflated, with rails 5 defining a heart receiving recess 6 in the central portion of the posterior surface. The balloon is sealed to the tube at the distal end 7 of the balloon and the 5 proximal end 8 of the balloon. The distal end of the tube may extend a short distance from the distal end of the balloon, and is fitted with an atraumatic tip 9 which can help dissect tissue when the device is pushed into the chest cavity. The portion of the rigid tube inside the balloon has inflation outlets 10, seen 10 in Figure 5, which allow inflation

fluid to pass into and out of the balloon. Referring again to Figure 1, near the proximal end 11 of the rigid tube, a handle 12 is fitted to the tube and a fluid tight connector 13 such as a typical Luer fitting, quick disconnect or other type fitting 15 is fitted to the tube in order to provide a fluid path from the air pump, through the rigid tube and into the balloon.

Figure 2 shows another shape for the cardiac pump. The balloon has an annular bulge around its perimeter which defines a recess 6 in the center or the posterior surface 4 which will 20 fit over the heart and rest on the rounded anterior surface of the heart. The overall shape of the posterior surface 4 in both Figures 1 and 2 is gently cupped to receive the heart, and maybe described as pal mate or palm shaped, such that it conforms to the gently curved anterior surface of the heart. This shape 25 permits

placement of the balloon over the heart and effective inflation to compress the heart, while also assisting in maintaining the balloon in place over the heart. A slightly convex shape on the posterior surface is also useful to compression of the heart. 5 30 Merely spherical or bulb shaped balloons may also be used.

Figure 3 shows an optional plastic introducer sleeve 14 with a rounded distal tip which may be used to facilitate insertion of the balloon into the chest. Each lateral corner 20 (indicated in Figure 6) of the balloon is rolled toward the center of the balloon, and the device is slipped into the introducer sleeve. The sleeve is a tear away sheath and has scored line 15 on the side to allow the sleeve to be removed from the device by tearing the sleeve along the scored line. The scored line is provided along the side of the device so that side-ways pull is necessary, thus avoiding the need for any upward force which could cause the tip to be forced downward into the heart, a concern that will become apparent from the description of use provided below. The pull tab 16 also serves as a distinguishing shape indicating the front of the device, and serves as a physical stop which prevents incorrect insertion of the device (i.e. insertion with the anterior surface of the balloon facing the heart). Figure 4 shows the detail of the balloon 2 folded within the sleeve as viewed from the distal end of the device.

Figure 5 shows the preferred structure of the rigid tube 3. The tube has a bend 17 of approximately 15 degrees near the midpoint of the balloon 2. This bend helps orient the tube so that the tip is urged upward into contact with the sternum rather than into softer tissue in the thoracic cavity, while allowing the device to be pushed upward into the thoracic cavity at a natural angle. The bend may be any convenient angle which facilitates insertion. The inflation ports 10 are shown along the distal portion of the rigid tube, corresponding to that part of the tube surrounded by the balloon. The atraumatic olive tip 9 prevents

unnecessary tearing while allowing for any needed dissection of tissue away from the sternum to permit passage of the balloon device into the substernal space above the heart. The proximal section of the tube 11 is provided with any type of quick 5 disconnect fitting 13 typically used for medical devices. The handle 12 shown in Figures 1 and 5 may be made in any comfortable size, or omitted altogether. The top or anterior surface of the device, which is the preferred orientation for insertion and use, may be marked on the tube or handle in any 10 suitable manner, either with printed instructions, color coding, texture or shape. The anterior side of the tube may be fitted with large fin which does not permit incorrect insertion. Also, the length of the inserted portion of the balloon device can be indicated by markings, such as ruler markings on the rigid tube, 15 in order to provide medics with an indication of how far the device is inserted, thus ensuring its proper placement in position over the heart. The rigid tube may be made of any material sufficiently rigid to permit insertion into the thoracic cavity through the prepared channel. Stainless steel, 20 nitinol and other metals, hard plastics, graphite composites and numerous other materials may be used.

The balloon 2 shown as shown in Figure 1 is fabricated of pieces as shown in Figures 6a, 6, 6c and 7. The balloon top piece 19 shown in Figure 7 is made of a single piece of plastic. 25 The series of steps use to create pleats which, when fill with air comprise the rails 5 of the balloon shown in Figure 1, is illustrated in Figures 6a, 6b and 6c. In Figure 6a, the plastic sheet 20 is folded along the crease 21 and the folded portion 22 is heat sealed together with the major portion along heat seal 30 line 23. The waste area 24 is trimmed away and the top piece appears as illustrated in Figure 6b with pleat 25 formed. The

top piece, when unfolded and laid flat, appears as shown in Figure 6c, with the pleat 25 extending upwardly from the sheet. The process is repeated on the other side of the sheet 20 to form a second pleat 26. When the top piece is complete, as 5 illustrated in Figure 7, it is heat sealed to the balloon bottom piece 27 along generally diamond shaped outline 28 and sealing darts 29 shown in dotted lines. The overall diamond shape has rounded corners 30 on the lateral corners of the balloon and extended corners 31 on the distal and proximal ends of the 10 balloon. These extended corners are used to seal the balloon to the insertion tube.

The cross section of the balloon in its deflated state is shown in Figure 8, while the cross section of the balloon when inflated is shown in Figure 9. When inflated, the pleats 25 and 15 26 form inflated rails 32 and 33 extending parallel to the center axis of the balloon, with a recess 34 between the rails, so that the balloon, when properly place over the heart, will curve around the surface of the heart and partially enclose the heart. The rigid tube 3 fits between the sealing darts 30.

20 The balloon of Figure 2 may be formed of pieces as shown in Figure 10. The posterior surface 4 is made from a single sheet of plastic 20 in the same manner as that described above in reference to Figures 6a, 6b, 6c and 7. for the posterior surface of the balloon of Figure 1. The sheet 20 is provided with pleats 25 35 and 36, which are formed in the same manner as illustrated in Figures 6a, 6b, and 6c. Heat seal lines 37 have already been heat sealed together. Excess material has been trimmed from pleats 35 and 36. Perpendicular pleats are then formed by folding the sheet along dotted lines 38 and heat sealing the 25 30 seam together along heat seal lines 39 and trimming away excess

plastic. Four pleated tabs will remain after excess plastic above the heat seal lines is removed. The bottom piece 41 is aligned with the top piece and the two are sealed together around outline 42 and the diamond shaped balloon with a ring of 5 pleats on the posterior surface is formed. When inflated, this balloon will appear as illustrated in Figure 2.

The balloon pieces are made of plastic such as polyurethane, polypropylene or polyethylene. The pieces are joined by heat sealing or adhesive. Balloon wall thickness of a 10 few mils (4 mils) is sufficient to ensure adequate strength of the balloon. The balloon may also be made of elastomeric materials, rubber, thermoplastic elastomers, latex, etc., and other non-elastomeric materials such as PTFE, canvas, woven polyester, nylon or sailcloth, etc.. After joining the pieces 15 together, the extended corners 21 form sleeves. The rigid tube is slipped into the balloon through the sleeves, and the sleeves are sealed to the tube to form an airtight bladder mounted on the rigid tube. For insertion the rounded corners of the balloon may be rolled inward toward the rigid tube, as shown in Figure 20 4, and covered with a tear-away sheath, an introducer sheath, or a thin walled cannula which is pulled back after the balloon is in place.

It will be appreciated that more rudimentary embodiments and more complex embodiments of the balloon will also be useful. 25 A simple pillow shape, which may be accomplished by heat sealing two diamond shapes together, or, for a fatter balloon, may be made in accordance with the steps described but without sealing the darts 29 to create the recess 6 shown in Figures 1 and 2. A mere round balloon mounted on a cannula or catheter can be used 30 to take advantage of the method presented herein while producing

some beneficial level of heart compression, and very complex geometries may be envisioned to distribute inflation pressure more precisely and evenly about the heart and produce optimal levels of heart compression. It should also be appreciated that 5 the rigid tube used for inflation may be omitted in favor of more complex insertion mechanisms. The balloon may be rolled inside the lumen of a cannula, catheter or trocar and pushed from the lumen after insertion of the cannula, catheter or trocar into the chest. Inflation and deflation may then be 10 accomplished by a flexible tube attached to the balloon.

Figure 11 shows the insertion point of the cardiac pump. The rib cage is comprised of the ribs 43, sternum 44, the costal cartilage 45 which joins the ribs to the sternum, and the xiphoid process 46. The diaphragm 47 separates the thoracic 15 cavity from the abdomen. The heart 48 is located in the center of the chest under the sternum, just above the diaphragm. A small vertical incision 49, just about 15 mm in length, is made just below the sternum 44 and the xiphoid process 46.

Figure 12 shows a sagittal cross section of the human chest 20 showing proper insertion pathway for placement of the cardiac pump into the chest of a cardiac arrest victim. The area delineated by the dotted line 50 is called the mediastinum. The heart is located in the middle mediastinum and the balloon is to be inserted into the anterior mediastinum. This space, 25 indicated by arrow 51, is also referred to as the substernal space or the sternocostal space. The medic uses an index finger 52 to push a hole up under the sternum 44 and through the diaphragm 47 and into the anterior mediastinum or sternocostal space 51 between the sternum and the heart. A blunt dissection 30 tool may be required, and mayo scissors may be used to help

punch the hole through the diaphragm. Thus, the cardiac pump may be placed, as illustrated in Figures 11 and 12, without substantially altering, disrupting or cutting through the rib cage, and without need to cut or retract bone or cartilage, and 5 incision through soft tissue is all that is required to insert the pump into the substernal space.

Figure 13 shows a cross section of the human chest showing the placement of the cardiac pump. The tip of the cardiac pump 1 is inserted into the small incision and pushed along the channel 10 formed by the finger, until it pushes through the hole in the diaphragm 47 and rests in the anterior mediastinum (or substernal or sternocostal) space 51 between the heart and the sternum. The 15 degree bend in the rigid tube permits easy insertion of the device while urging the distal tip upward 15 toward the sternum and away from the heart. The ideal position for the cardiac pump is directly above the heart. Note that the heart is surrounded by the pericardial sac 54 and that the pericardial sac need not be penetrated for operation of the cardiac pump.

20 When in place, the balloon portion 2 of the cardiac pump is inflated, as illustrated in Figure 14. The balloon expands within the chest, against the sternum 44 above and the heart 48 below. The sternum provides a rigid and immovable support for the balloon, so that the balloon expands toward the heart and 25 squeezes and compresses the heart. The heart is confined on the posterior side by the aorta, spinal column and other structures. Compression of the heart causes the heart to pump blood, as in the normal systole. After inflation has compressed the heart, the balloon is deflated and the heart relaxes and expands, just 30 as it does in the normal diastole and draws blood from the

pulmonary vein (not shown) as it does during normal operation. Upon deflation, the rounded corners 20 of the balloon will remain in place over the heart, as shown in the deflated position illustrated in Figure 15. The anterior view of the 5 balloon, corresponding to the Figure 14 and 15, is shown in Figure 16. The balloon 2 substantially covers the heart 48 in the anterior view.

The insertion point for the balloon, just below the point of the xiphoid process, is preferred because, as illustrated, 10 the pathway into the sternocostal space leads directly over the heart without encountering any other significant organs. Thus the danger of damage to other organs, such as the lungs, aorta, spinal column, etc. is minimized. If required, the device can be inserted through an incision in the intercostal spaces (spaces 15 between the ribs) 55 or even from an incision near the jugular notch 56 or clavicular notch 57 at the top of the sternum, indicated in Figure 11. Insertion through these alternative pathways will require sliding the cardiac pump near the lungs or brachio-cephalic vessels, and extra care must be exercised to 20 avoid damage to these structures.

The method of implanting the cardiac pump described above does not require disruption or gross invasion of the rib cage which is required for open heart massage. There is no need to cut the bone or cartilage of the rib cage, and no need to 25 retract the rib cage into an open position to gain access to the heart. This permits direct cardiac massage immediately upon cardiac arrest. The device can be inserted in non-sterile environments by paramedics as soon as they reach a cardiac arrest victim. If a cardiac arrest victim has recently undergone 30 open chest surgery, use of the device allows direct cardiac

massage without the need to disrupt the recently closed rib cage, and eliminates the danger of closed chest compression on the healing rib cage. It may even prove beneficial to implant the cardiac pump immediately after open chest surgery for 5 patients at high risk of perioperative cardiac arrest (thus the utility of an embodiment constructed without the rigid tube which facilitates insertion). In this case, the balloon may be secured in place with resorbable sutures, suturing the balloon to the anterior chest wall or rib cage.

10 The mechanism for continuously inflating and deflating the cardiac pump may be provided in a variety of embodiments. Shown in Figure 17 is one such system, comprised of a standard dual acting pump 58 such as the Guzzler™ 1.5 inch hand pump available from The Bosworth Company of East Providence, RI. The pump is 15 hand operated, and may also be operated by motor 59 through releasable linkage or clutch 60. On each down-stroke, a certain amount of air is forced into the cardiac pump balloon 2, and on each up-stroke, the air is drawn from the cardiac pump balloon. The specifications for the exact amount of air should be chosen 20 to avoid the possibility of bursting the balloon and also to ensure adequate inflation pressure for compressing the heart. The positive displacement pump with an output matched to the volume of the expanded balloon is preferred. The air pump 58 is attached to the cardiac pump 1 through a supply line 61 of tygon 25 tubing or other flexible tubing hooked up to the connector 13 on the cardiac pump 1. A pressure relief valve 62 can be provided on the supply line, or at any convenient point in the system. A low pressure warning sensor can be provided in the supply line or at any convenient point in the system, its purpose being to 30 indicate a low pressure condition which would indicate the balloon has ruptured or that the air supply line to the cardiac

pump is leaking (and, consequently, the cardiac pump is not inflating to compress the heart and provide the needed cardiac massage). The pump handle may be attached to the small electric motor and operated automatically, rather than manually, thus

5 eliminating the fatigue of extended operation by hand. The motor is operated by battery or household current adapter or power supplies available in ambulances. In case of failure of the electric motor, the pump handle may be uncoupled from the motor allowing the pump to be operated by hand.

10 Use of an automatic inflation system is preferred in order to provide consistent cardiac massage over an extended period of time. The speed of the pump can be adjusted to achieve the desired pace of compressions per minute. Currently, it is standard to provide 80 compressions per minute, but some studies

15 indicate that 120 compressions per minute in a technique called high impulse CPR and active compression/decompression may be better. An adjustable rate of compression will provide for optimization of the compression rate. The amount of air provided in each stroke may be set by choosing a positive displacement

20 pump with an appropriate output volume or by regulating the pressure output of the pump. The stroke of the pump or the pressure limit may be adjusted to achieve optimal compression. In this manner, the rate and the amount of inflation can be provided more consistently with the cardiac pump when operated

25 by motor than when operated by hand, and the problem of operator fatigue is eliminated.

Inflation air may be provided by other means, such as a centrifugal air pump or compressor or compressed air tank. Where continuous air supply is used, such as with a centrifugal air pump, the cycling of the cardiac pump may be accomplished by

venting air from the supply line and the balloon at regular intervals, thus allowing the natural resiliency of the thoracic cavity and the heart to act upon the unpressurized balloon to deflate the balloon and allow the heart to expand and take in 5 blood from the lungs. The high pressure source used to inflate the balloon can also, through the use of a Venturi vacuum generator, be used to actively deflate the balloon. For active deflation of the balloon while using a high pressure gas source, the pressurized air used to fill the balloon can be alternately 10 bypassed (using a three way valve) to a Venturi vacuum pump which draws suction on the balloon. The alternate cycling can be accomplished with a solenoid operated valve, a pneumatically operated valve, or manually, with appropriate valve configurations for aligning gas flow for inflating and deflating 15 the balloon. While air has been suggested as the preferred inflation fluid, water, saline solution, CO₂, inert gases, etc. may be substituted as the inflation fluid.

It will at times be desirable to provide for defibrillation of a heart attack victim while the balloon is in place. This may 20 happen, for example, when a heartbeat is restored with treatment after a period of cardiac massage but fibrillation occurs for which the indicated treatment is electrical cardioversion rather than cardiac massage. Thus a balloon that is combined with defibrillation electrodes is desirable. Figures 18 and 19 show 25 an embodiment of the cardiac pump 1 which is fitted with several patch electrodes 63 on the posterior surface 4 of the balloon 2. (Note that the balloon in this embodiment is another embodiment of balloon shape, and resembles a simple pillow shape.) The electrodes are connected to a standard defibrillator power 30 source through insulated electrical wires 64 which are potted to

the electrodes with a flexible electrical epoxy and passed through cable sheath 65 attached to the insertion rod 3.

When the balloon is in place within the patient, in the anterior mediastinum, the posterior surface with the electrodes 5 will face the heart, and be in contact with the anterior surface of the pericardium. Electrical activity of the heart can be monitored through the pericardium, and the electrical stimulus required for cardioversion can be applied through the pericardium. The electrodes may be used as dual purpose sensing 10 electrodes and electrical stimulus electrodes, or they may be provided as a pair of sensing electrodes 63a and 63e and a pair of cardioversion electrodes 63b and 63d. The electrodes may be used as monopolar electrodes or bipolar electrodes, as medically indicated. The electrodes are made of standard material for 15 patch electrodes and surface electrodes, and may also be made of simple electrically conductive sheaths of, for example, copper or other metal or electrically conductive polymers. The electrodes are attached to the balloon with flexible adhesive. The electrical wires 64 are secured to the balloon with adhesive 20 or an overlying layer of balloon material heat sealed to the balloon over the wires.

A sensing circuit may be included in the defibrillation power source circuitry for the purpose of monitoring the heartbeat when restored. The sensing circuit and monitor can be 25 used to alert the medic that defibrillation has occurred. Cardioversion can be accomplished automatically upon sensing fibrillation or absence of heartbeat, as is done with pacemakers, or may be accomplished manually through the balloon mounted electrodes. The sensor and alarm circuitry may be 30 interconnected with the air pump control circuitry in order to

stop the cardiac pump during defibrillation. The air pump may be stopped with the balloon inflated or deflated to ensure optimum contact with the pericardium during delivery of electrical stimulus through the electrodes.

5 All of the component parts of the device can fit into a kit the size of a brief case or suit case, and can be made readily transportable for use in any environment. In an embodiment intended for paramedic and ambulance use, it is expected that the motor, pump and an electric battery will be provided in
10 single case which also holds several cardiac massage balloon assemblies and several lengths of supply line to permit use in any situation. In an embodiment intended for use in hospitals and emergency rooms, the components of the entire system may be provided in a rolling cart, and may be included in crash carts.

15 The repeated inflation and deflation of the cardiac pump can be continued indefinitely, and can be stopped when the cardiac arrest victim is resuscitated and recovers a normal heartbeat, or until the victim is ready for reparative surgery, such as a coronary bypass, pacemaker implantation or other
20 surgery, or until it is determined that the victim cannot be revived. If the victim recovers a heartbeat spontaneously while the cardiac pump is operating, the pump may be stopped and the balloon deflated. This will allow the heart to beat without substantial interference from the balloon, but will leave the
25 balloon in place in case the victim's heart stops beating again. When the balloon must be removed, it is deflated and pulled out of the chest, and the flexibility of the balloon will allow easy removal through the small incision used for insertion. As part of the recovery process, the small incision in the skin and the

small puncture in the diaphragm may be closed with sutures or staples.

One way in which the use of the cardiac pump may be optimized is to use it in conjunction with a CO₂ monitor placed 5 in the victim's airway. The CO₂ content of the exhalant during CPR indicates the success of the resuscitation effort, according to Gudipati, et al., *Expired Carbon Dioxide: A Non-Invasive Monitor Of Cardiopulmonary Resuscitation*, 77 Circulation 234 (1988). By measuring the CO₂ in the expired air, medics can 10 determine whether or not the cardiac pump is performing adequately. If expired CO₂ measurements indicate that blood flow is insufficient, the volume and/or pressure of air injected into the cardiac pump balloon may be increased to cause more rapid expansion and larger expansion, or the pace of inflation and 15 deflation may be adjusted within the medically accepted ranges (currently 80-120 compressions per minute).

A wide variety of materials and components may be used to construct the cardiac massage balloon and the manual and motorized inflation systems described above. The materials and 20 specific components mentioned in connection with the devices have proven suitable for experimental use. It is specifically intended that balloon materials and configurations may be optimized upon further development and experience with the cardiac massage pump. Thus, while the preferred embodiments of 25 the devices and methods have been described, they are merely illustrative of the principles of the invention. Other embodiments and configurations may be devised without departing from the spirit of the inventions and the scope of the appended claims.

We claim:

1. A minimally invasive device for cardiac massage within the chest of a human, said device comprising:

5 a rigid insertion rod having a distal end and a proximal end, and having a fluid conduit communicating from the proximal end to the distal end, and having an fluid port in the distal end and a fluid port in the proximal end, said distal and proximal fluid ports communicating with the fluid conduit so that the fluid ports and fluid conduit provide a pathway for fluid to pass through the 10 rigid tube;

15 a balloon mounted on the distal end of the rigid insertion rod over the distal fluid port so that the balloon may be inflated and deflated by passage of fluid through the fluid conduit, said balloon and distal end of the rod forming an insertion portion;

20 said insertion portion being sized and dimensioned to enable insertion of the insertion portion within the chest of the human between the heart and the sternum of the human, when the balloon is deflated; and

25 said balloon having an inflated shape sized and dimensioned to compress the heart when said balloon is inflated within the chest of the human.

2. A minimally invasive device for cardiac massage within 25 the chest of a human, said device comprising:

30 a rigid insertion rod having a distal end and a proximal end, having a fluid conduit communicating from the proximal end to the distal end, and having an fluid port in the distal end and a fluid port in the proximal end, said distal and proximal fluid ports communicating with the fluid conduit so that the fluid ports and fluid

conduits provide a pathway for fluid to pass through the rigid tube;

5 a balloon mounted on the distal end of the rigid insertion rod over the distal fluid port so that the balloon may be inflated and deflated by passage of fluid through the fluid conduit, said balloon and distal end of the rod forming an insertion portion;

10 said insertion portion being sized and dimensioned to enable insertion of the insertion portion within the chest of the human between the heart and the sternum of the human, when the balloon is deflated;

15 said balloon having an inflated shape sized and dimensioned to compress the heart when said balloon is inflated within the chest of the human;

20 said rigid insertion rod having a bend therein of approximately 15° from the plane of the balloon;

a fluid connector on the proximal end of the rigid tube;

a fluid supply line connected to the fluid connector on the proximal end of the rigid tube; and

25 a fluid supply pump connected to the balloon through the fluid supply line, the fluid connector and the rigid tube and the distal fluid port in the rigid tube.

3. A method of performing direct cardiac massage within the closed thoracic cavity of a human without having to cut through 30 the ribcage, said method comprising:

providing an inflatable balloon sized and dimensioned to fit between the rib cage and the heart;

inserting the balloon into the thoracic cavity without disrupting the ribcage; and

5 inflating the balloon while it is in place within the thoracic cavity to cause compression of the heart.

4. The method of claim 3 further comprising the steps of repeatedly deflating the balloon to allow the heart to expand and inflating the balloon to compress the heart.

10 5. A method of compressing the heart with a balloon placed within the thoracic cavity of a human without need for disrupting the rib cage, said method comprising:

providing a balloon device for cardiac massage within the chest of a human, said device comprising:

15 a rigid insertion rod having a distal end and a proximal end, said rigid insertion rod having a fluid conduit communicating from the proximal end to the distal end thereof, and having a fluid port in the distal end and a fluid port in the proximal end, said distal and proximal fluid ports communicating with the fluid conduit so that the fluid ports and fluid conduits provide a pathway for fluid to pass through the rigid tube;

20 a balloon mounted on the distal end of the rigid insertion rod over the distal fluid port so that the balloon may be inflated and deflated by passage of fluid through the fluid conduit, said balloon and distal end of the rod forming an insertion portion;

25 said insertion portion being sized and dimensioned to enable insertion of the insertion portion within the

chest of the human between the heart and the sternum of the human, when the balloon is deflated;

5 said balloon having an inflated shape sized and dimensioned to compress the heart when said balloon is inflated within the chest of the human;

inserting the insertion portion of the balloon device into the thoracic cavity of the human, so that the balloon is situated between the heart and the sternum;

10 inflating the balloon while it is in the thoracic cavity, thereby compressing the heart as the balloon inflates.

6. The method of claim 5 further comprising the step of alternately deflating and inflating the balloon to cause repeated compression of the heart.

7. The method of claim 5 wherein the rib cage is not opened or 15 retracted away from the heart for the purpose of placing the balloon between the sternum and the heart.

8. The method of claim 5 wherein the balloon is inserted into the substernal space of the human.

9. The method of claim 5 wherein the balloon is inserted into 20 the anterior mediastinum.

10. The method of claim 5 wherein the balloon is inserted into the sternocostal space.

11. A cardiac pump system comprising:

25 an inflatable bladder sized and dimensioned to fit in the anterior mediastinum of a human;

pump means for repeatedly inflating and deflating the inflatable bladder while it is disposed within the anterior mediastinum of the human;

5 means for inserting the inflatable bladder into the sternocostal space of the human.

12. The device of claim 11 wherein:

10 the means for repeatedly inflating and deflating the inflatable bladder comprises a pump and an inflation fluid supply line, said fluid supply line connecting the pump to the inflatable bladder and allowing the passage of inflation fluid into the inflatable bladder.

13. The device of claim 11 further comprising:

a motor for energizing the pump means.

14. The device of claim 11 further comprising:

15 a motor and a portable power supply for said motor, said motor operably connected to the pump means to energize the pump means.

16. The device of claim 11 further comprising a pressure relief 20 means for limiting the pressure of inflation fluid in the inflatable bladder.

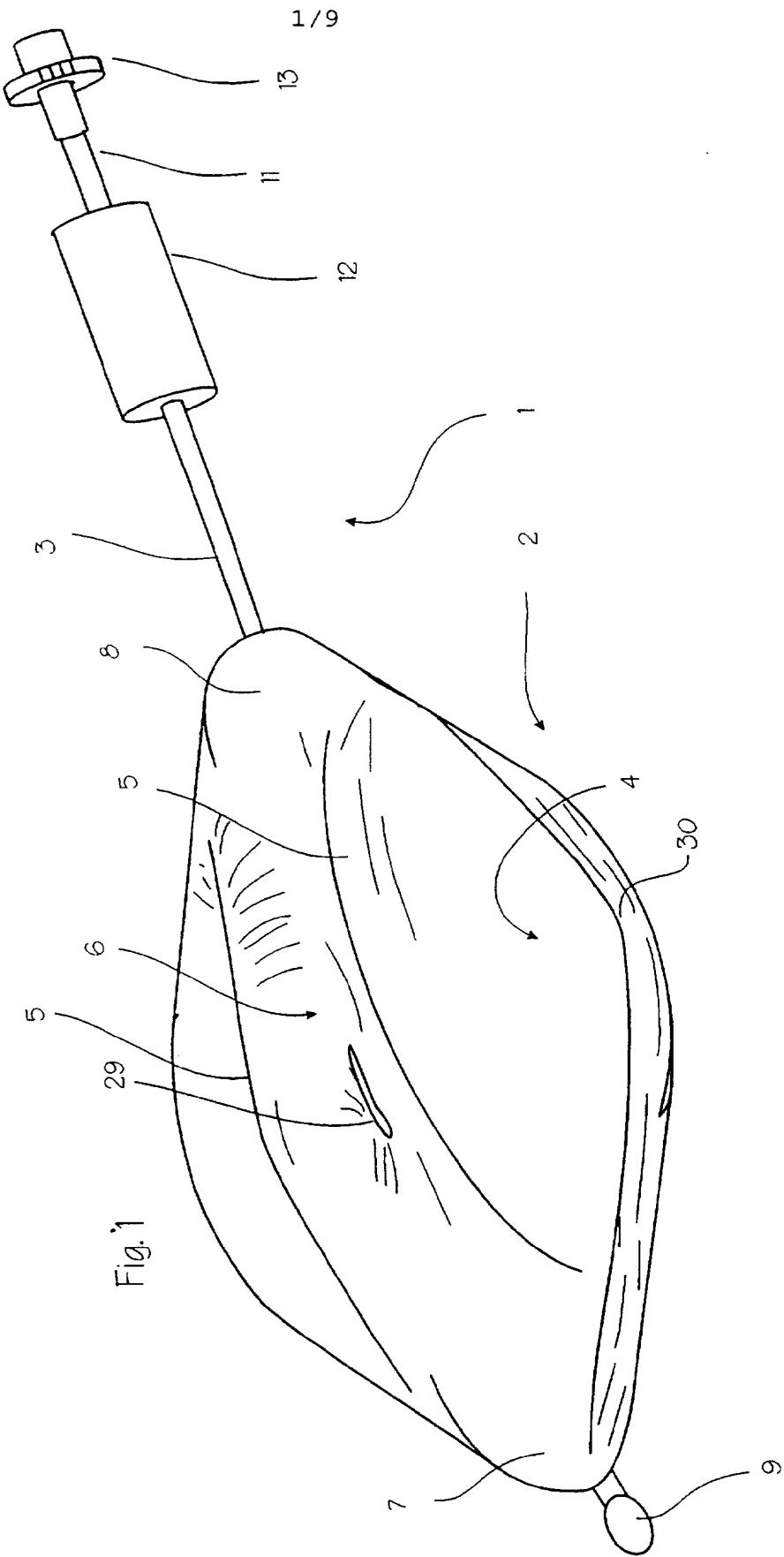
17. A diamond shaped bladder for insertion into the sternocostal space of a human, said bladder having an anterior surface and a posterior surface and a center axis running 25 through the upper and lower points of the diamond shaped bladder, the posterior surface of the diamond shaped bladder having rails which bulge posteriorly from the major portion of the bladder and are roughly parallel to the center axis.

17. The device of claim 1 wherein the balloon in its inflated shape is characterized by an anterior surface and a posterior surface, said posterior surface facing the heart when the balloon is in place and inflated between the sternum and the 5 heart, said posterior surface further characterized by a palmate shape.

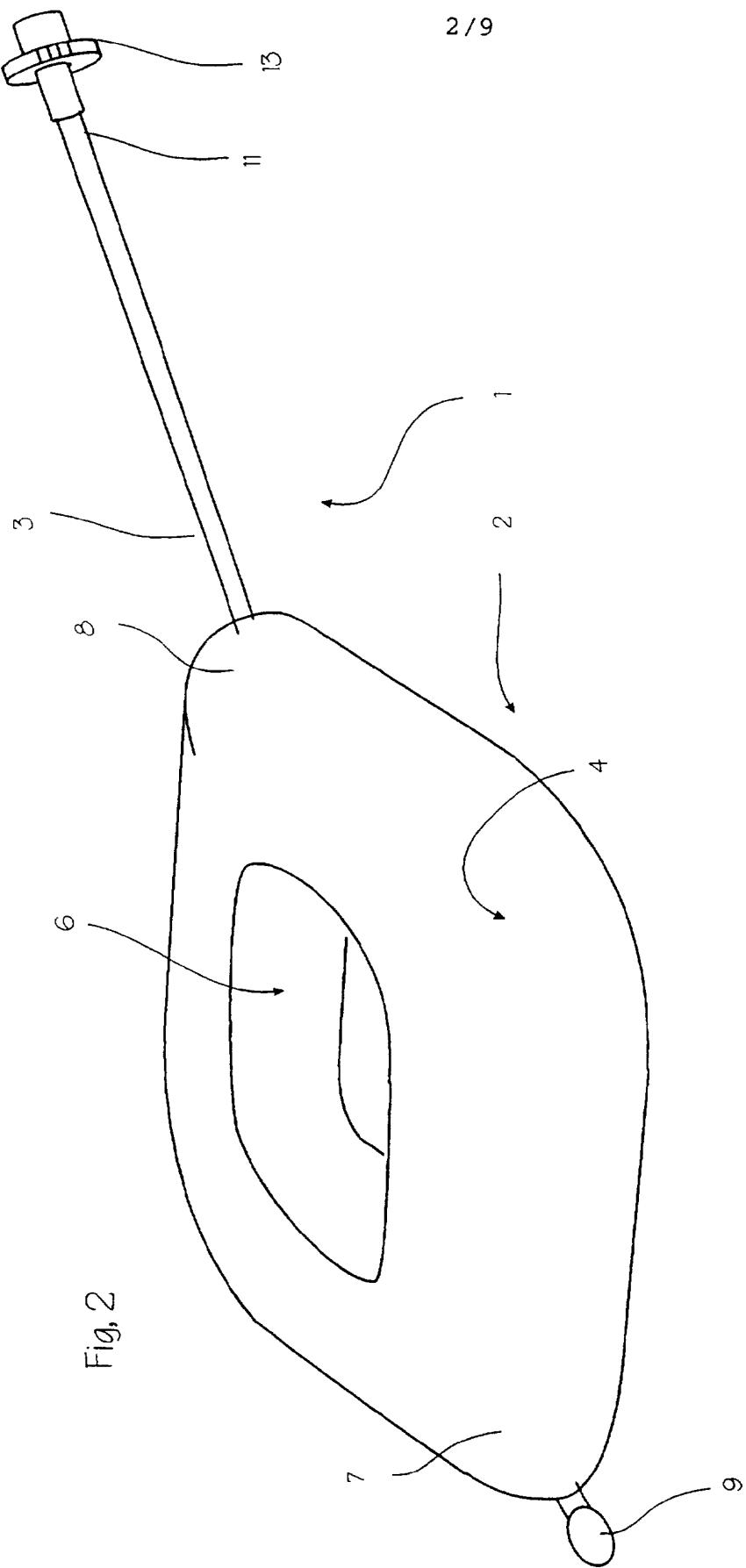
18. The device of claim 2 wherein the balloon in its inflated shape is characterized by an anterior surface and a posterior surface, said posterior surface facing the heart when the 10 balloon is in place and inflated between the sternum and the heart, said posterior surface further characterized by a palmate shape.

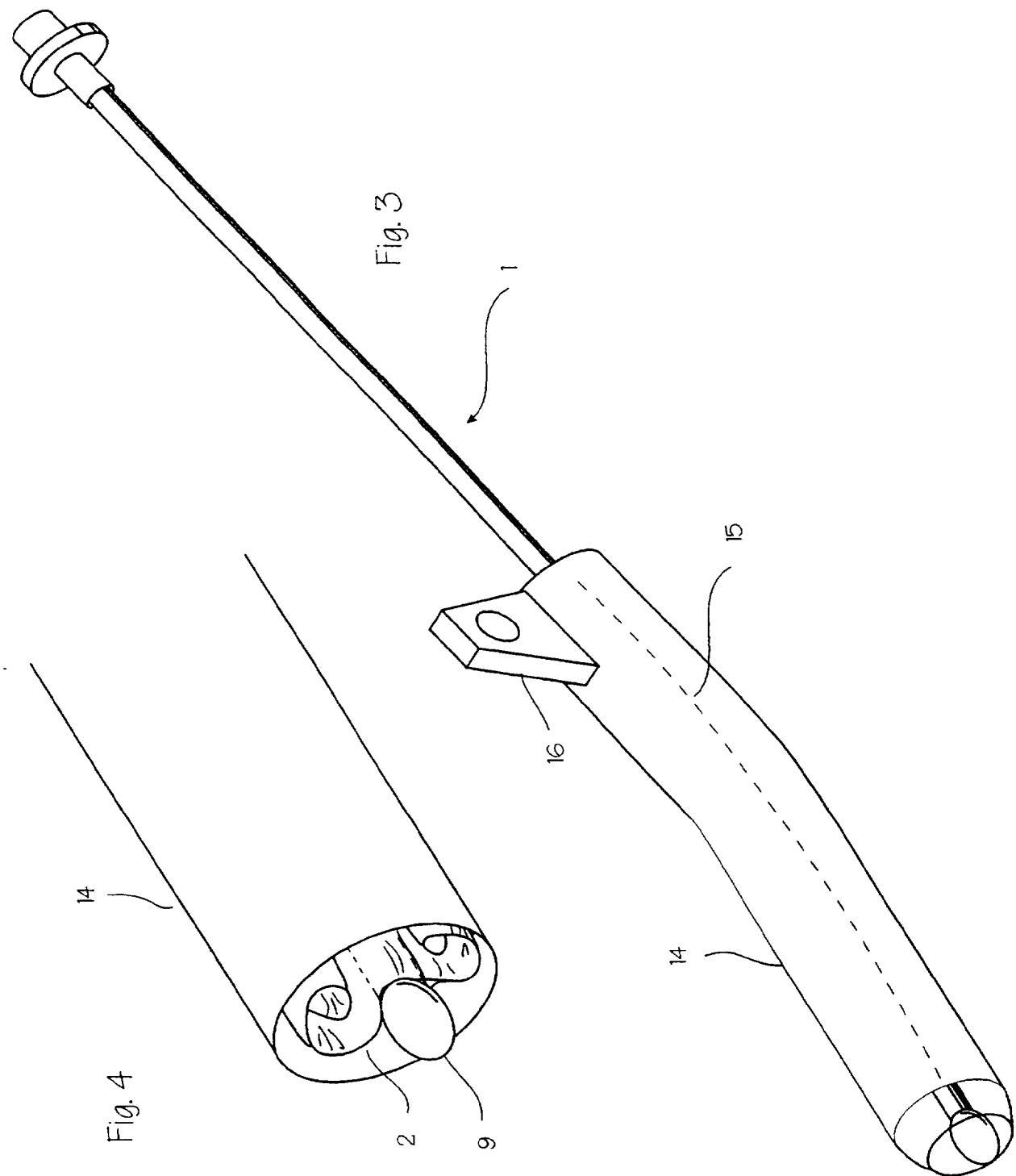
19. The device of claim 1 wherein the balloon in its inflated shape is characterized by an anterior surface and a posterior 15 surface, said posterior surface facing the heart when the balloon is in place and inflated between the sternum and the heart, said posterior surface further characterized by a convex shape.

20. The device of claim 2 wherein the balloon in its inflated 20 shape is characterized by an anterior surface and a posterior surface, said posterior surface facing the heart when the balloon is in place and inflated between the sternum and the heart, said posterior surface further characterized by a convex shape.



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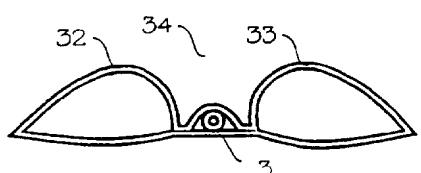
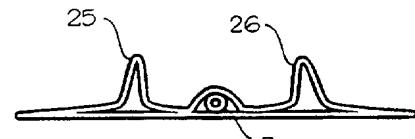
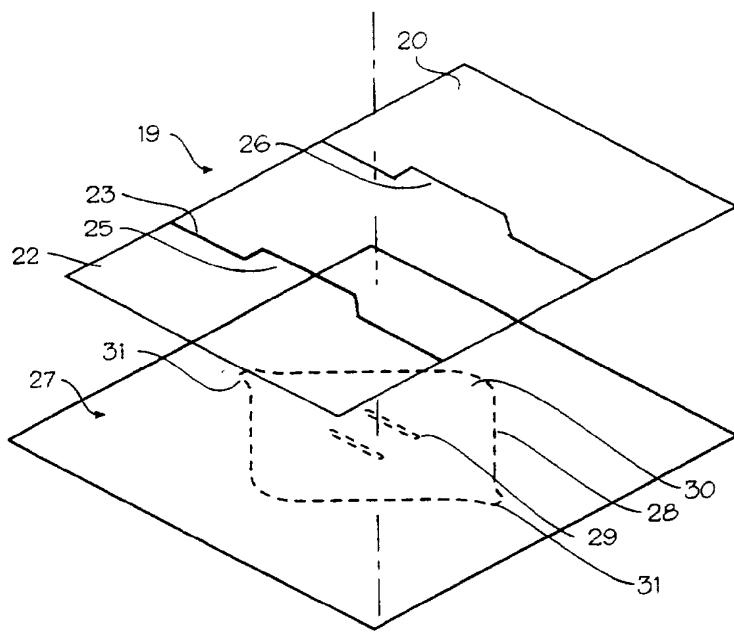
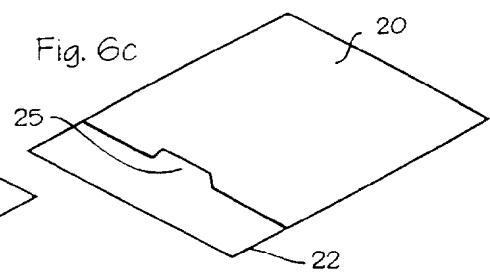
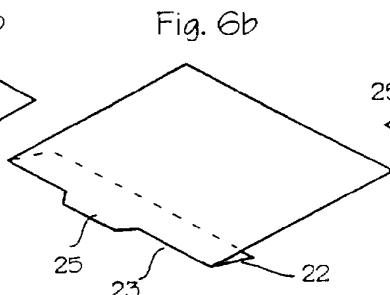
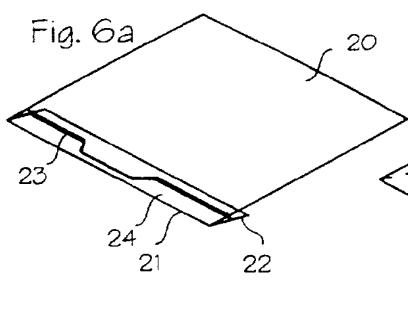
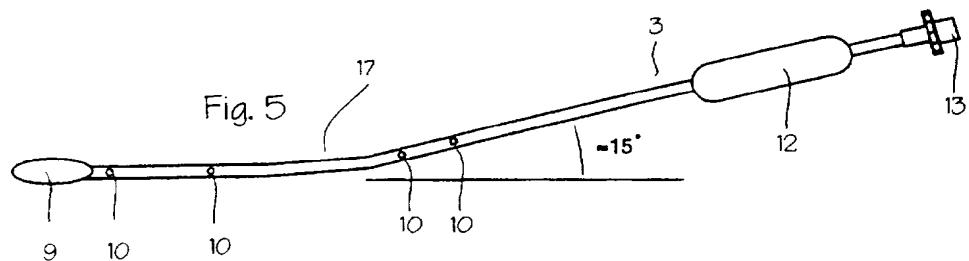
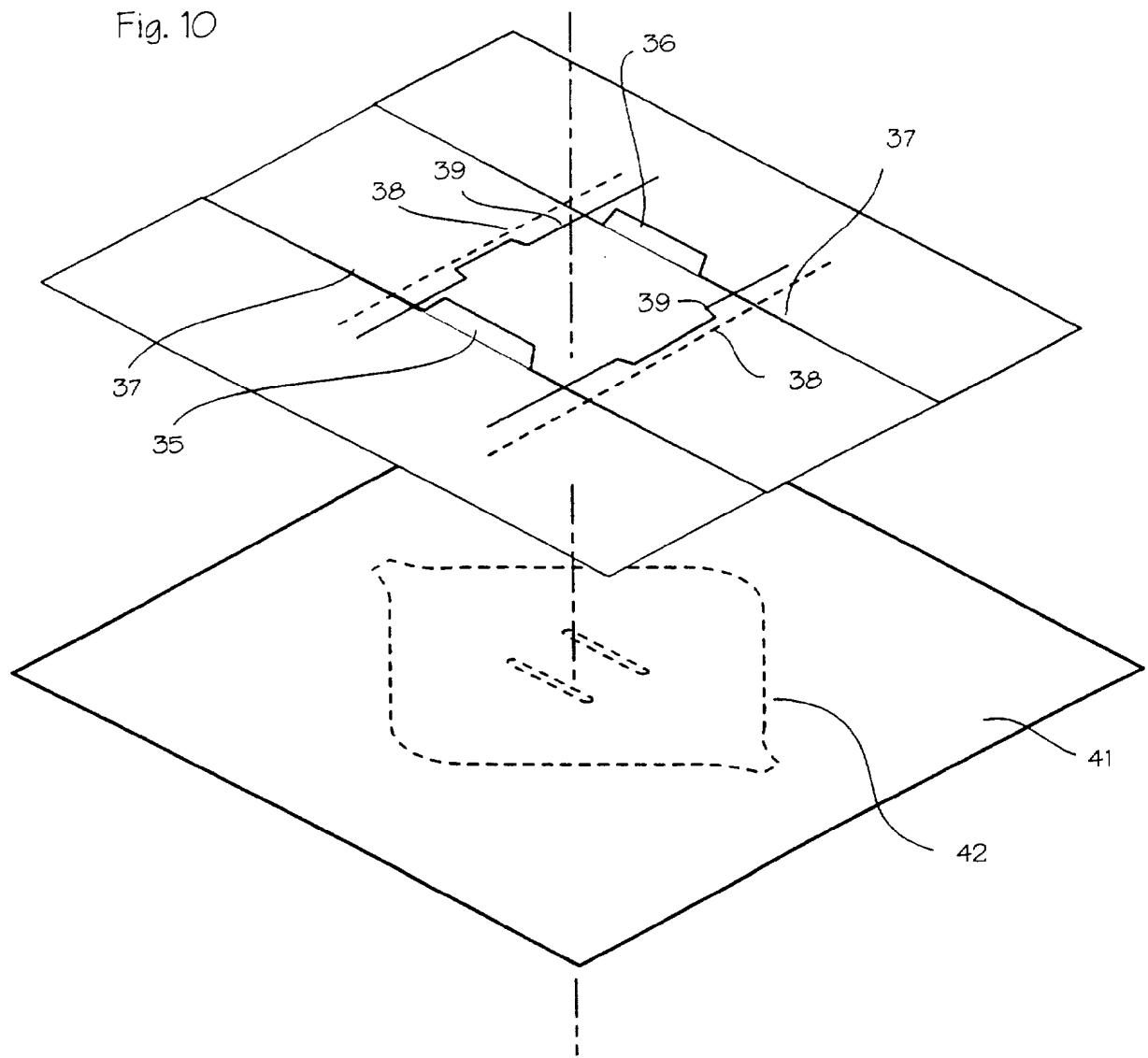
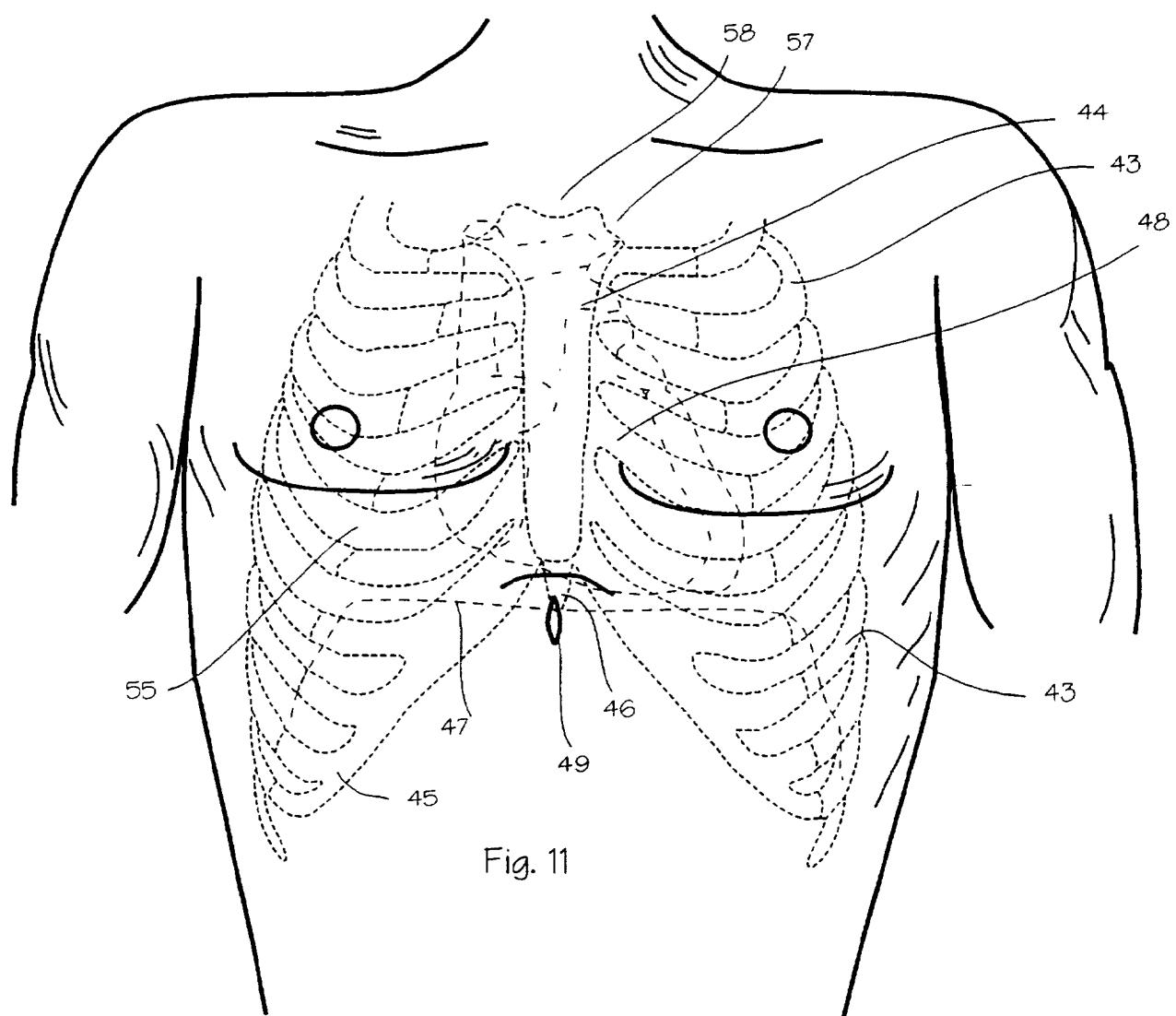


Fig. 10





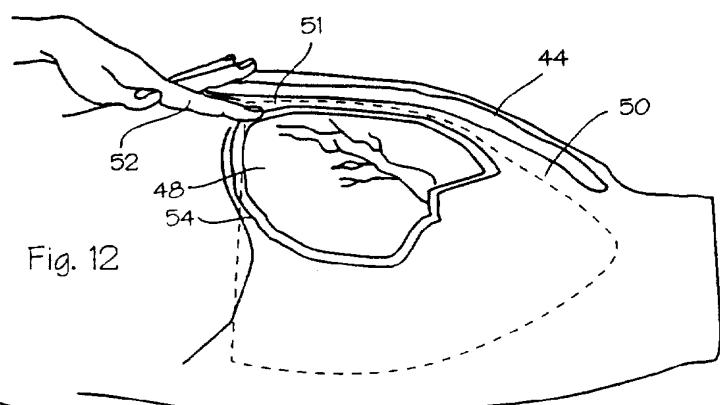


Fig. 12

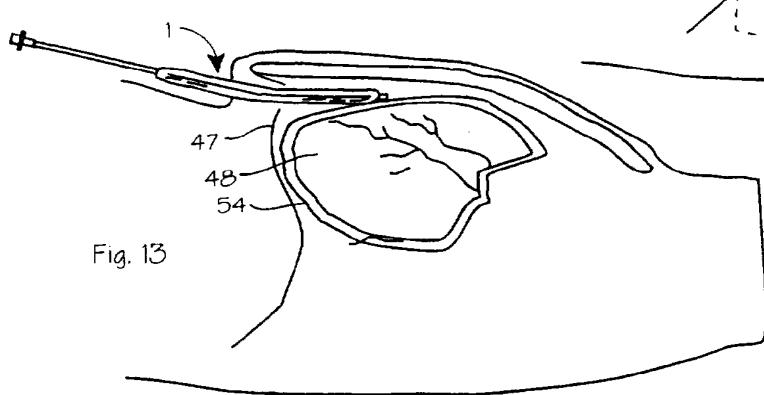


Fig. 13

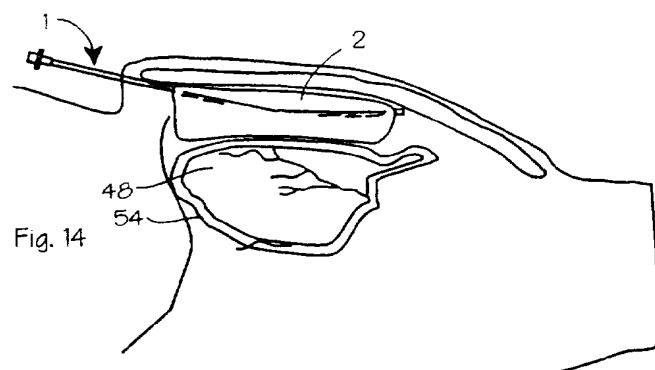


Fig. 14

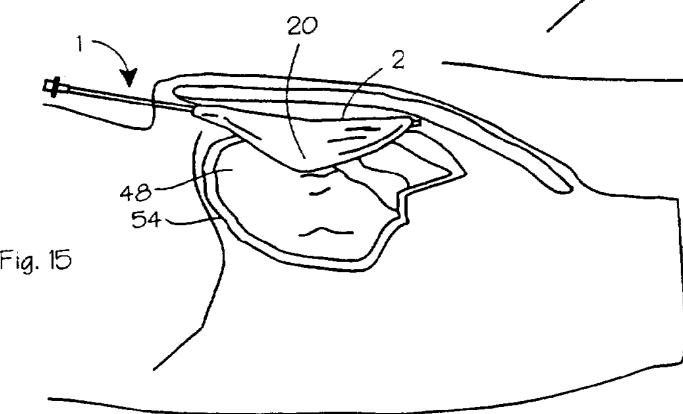


Fig. 15

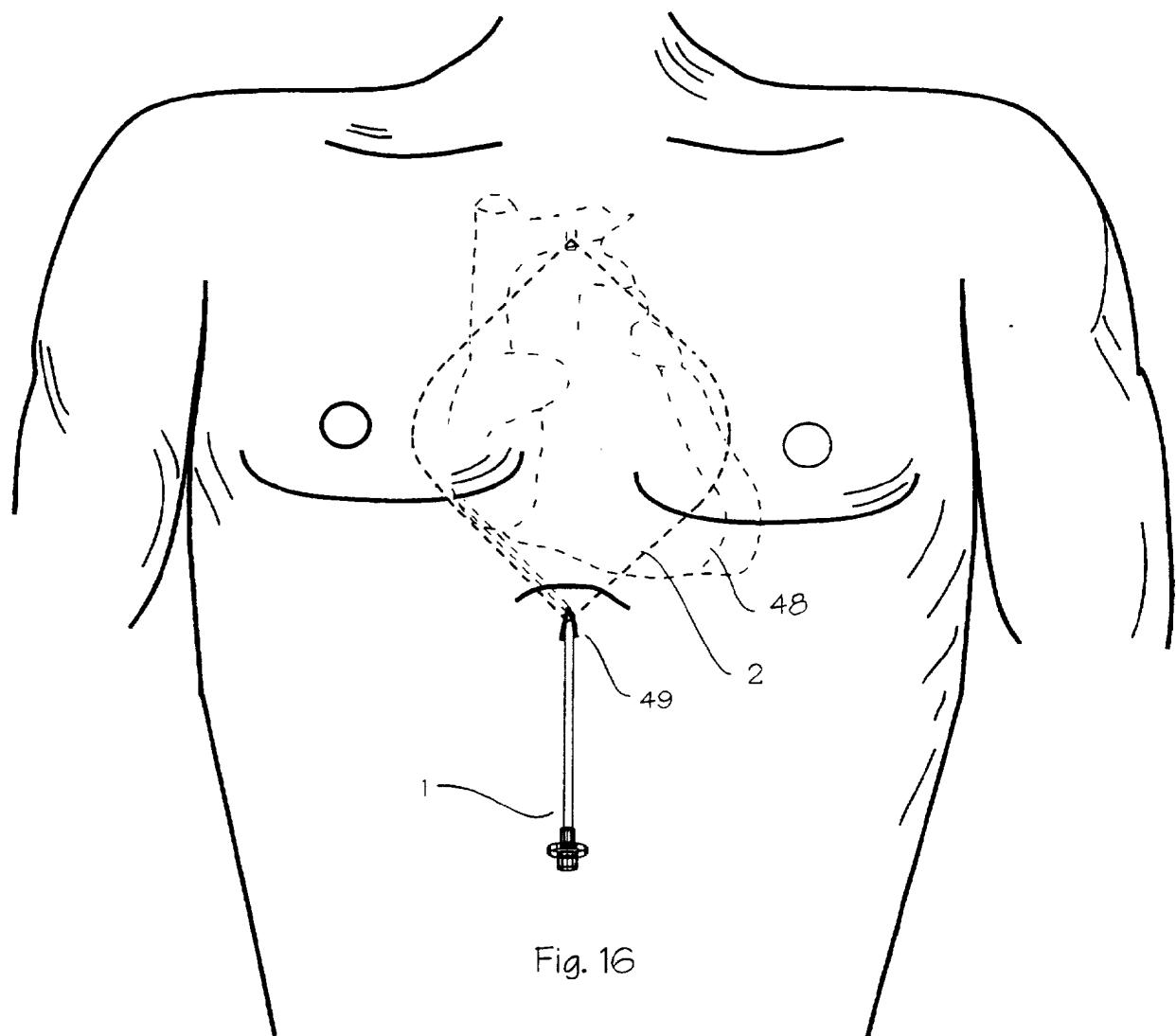
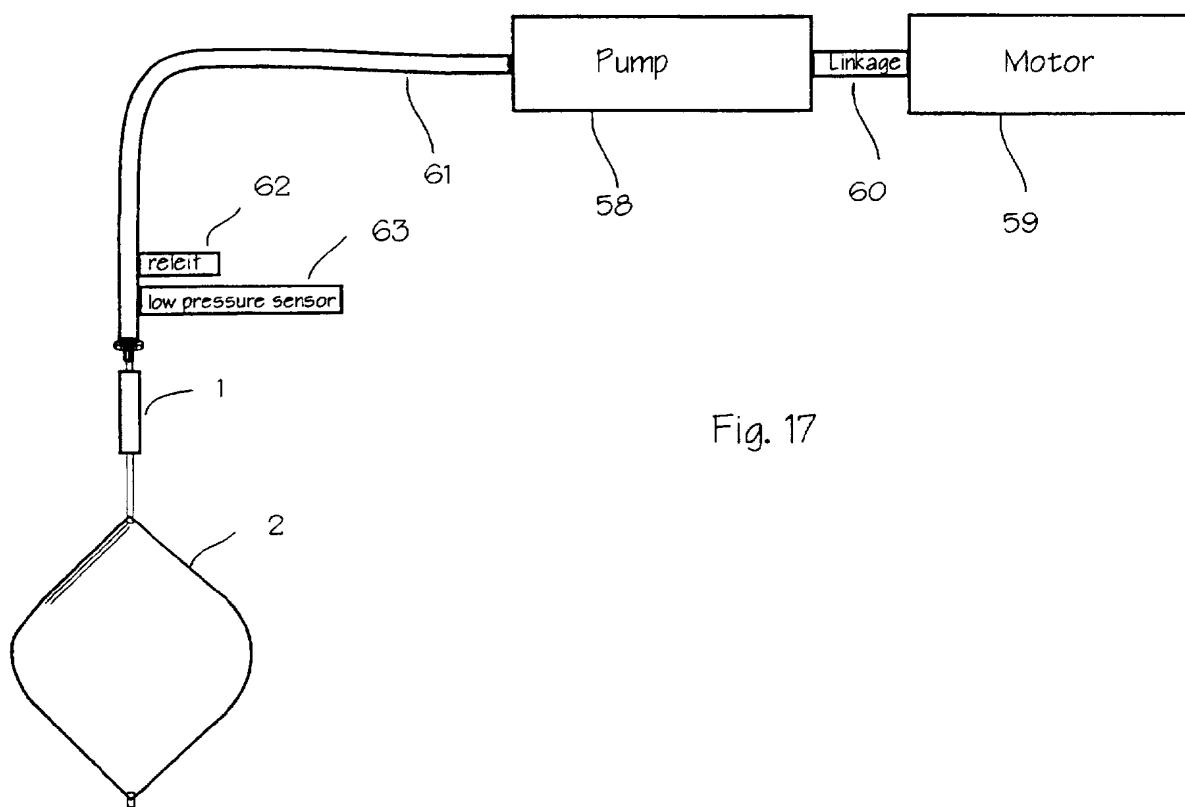


Fig. 16

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INTERNATIONAL SEARCH REPORT

International application No.
PCT/IB97/01096

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : A61H 31/00

US CL. : 601/153

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 601/151, 153

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 5,385,528 A (WILK) 31 JANUARY 1995. SEE ENTIRE DOCUMENT.	1-15
A		16-20

Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A"	document defining the general state of the art which is not considered to be of particular relevance	
"E"	earlier document published on or after the international filing date	"X"
"L"	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y"
"O"	document referring to an oral disclosure, use, exhibition or other means	"&"
"P"	document published prior to the international filing date but later than the priority date claimed	document member of the same patent family

Date of the actual completion of the international search

21 NOVEMBER 1997

Date of mailing of the international search report

08 DEC 1997

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